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PATENT

Appl. No. 10/658,823 Amdt. dated November 27, 2006 Reply to Office Action of July 25, 2006

## Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application:

## **Listing of Claims:**

1 1.-35. (Canceled) 36. (Previously Presented) A method of identifying an inhibitor of a 1 2 glycosyltransferase that transfers a monosaccharide from a sugar nucleotide to an acceptor 3 substrate, the method comprising 4 contacting the glycosyltransferase, an acceptor substrate, and a donor substrate with a hydrophobic, non-carbohydrate test compound that inhibits interaction of a sugar with 5 hydrophobic amino acids in the active site of the glycosyltransferase and 6 determining the degree to which the activity of the glycosyltransferase is inhibited 7 8 in the presence of the test compound. (Previously Presented) The method of claim 36, wherein the activity of 1 37. the glycosyltransferase is determined using an antibody that is specifically immunoreactive with 2 3 a product of the reaction catalyzed by the glycosyltransferase. (Previously Presented) The method of claim 37, which is an ELISA 1 38. 2 format. 39. (Previously Presented) The method of claim 36, wherein the 1 2 glycosyltransferase is expressed in a recombinant cell. 1 40. (Previously Presented) The method of claim 36, wherein the donor 2 substrate or acceptor substrate is labeled. 1 41. (Withdrawn) The method of claim 40, wherein the label is a radioactive 2 label.

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(Withdrawn) The method of claim 41, which is a radioactive column 42. 1 2 assay. (Previously Presented) The method of claim 40, wherein the label is a 43. 1 2 fluorescent label. (Previously Presented) The method of claim 36, wherein the 44. 1 glycosyltransferase is a fucosyltransferase. 2 (Withdrawn) The method claim 36, wherein the glycosyltransferase is a 1 45. 2 sialyltransferase. (Withdrawn) The method claim 36, wherein the glycosyltransferase is an 46. 1 2 N-acetylglucosaminyltransferase. (Canceled) The method of claim 36, wherein the compound comprises an 47. 1 aromatic or aliphatic ring structure. 2 (Withdrawn) The method of claim 36, wherein the compound comprises 48. 1 2 an aryl moiety. (Previously Presented) The method claim 36, wherein the compound 1 49. comprises a heteroaryl moiety. 2 (Previously Amended) The method of claim 49, wherein the heteroaryl 50. 1 moiety is selected from the group consisting of a thiophene, pyridine, isoxazole, phthalimide, 2 pyrazole, indole, quinoline, phenothiazine, carbazole, benzopyranone, and a furan group. 3 (New) The method of claim 36, wherein the hydrophobic, non-51. 1 carbohydrate test compound comprises a member selected from the group consisting of a 2 heteroaryl moiety having from 5 to 16 ring members wherein from 1 to 3 ring members are each 3 independently selected from the group consisting of N, O and S wherein the heteroaryl ring 4

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- structure is optionally substituted, and an aliphatic ring structure having from 3 to 7 ring 5
- members and is optionally substituted. 6